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Washington, D.C. 20231

Submitted herewith for filing is the patent
 application of:

John R. Fyson

For: METHOD OF TREATING WASTE
 EFFLUENT

Enclosed are:

- ☒ 6 sheet(s) of drawing(s).
- ☒ An assignment of the invention to Eastman Kodak Co.
- ☒ A certified copy of a application.
- ☐ An associate power of attorney.
- ☐ A Disclosure Statement Under 37 C.F.R. 1.97.
- ☒ Combined Declaration for Patent Application and Power of Attorney.
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Original

Application Based on

Docket 71442JRE

Inventors: John Richard Fyson

METHOD OF TREATING WASTE EFFLUENT

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METHOD OF TREATING WASTE EFFLUENT

Field of the Invention

The present invention relates to a method of treating waste effluent, in particular photographic effluent, which contain reduced species, especially sulphur-oxygen species, such as, for example, thiosulphate or sulphite. The present invention also embraces apparatus for performing such methods of treatment.

Background of the Invention

It is usual for photographic effluent for both black-and-white and color processing to contain one or more reduced sulphur-oxygen species. Thiosulphate or "hypo" is commonly used as a fixing agent, and sulphite ions are often included in developer solutions and/or stop/clearing baths as a preservative scavenger for oxidized developing agent. Sulphite is also known as a fixing agent. DE-A-3635219 to AGFA-GEVAERT AG discloses treating dilute aqueous solutions of thiosulphate by oxidation with hydrogen peroxide in the presence of a chromium, vanadium, tungsten or molybdenum compound as catalyst and discloses that molybdate is preferred. It is desirable to oxidize thiosulphate in photographic effluent before discharging the effluents to a sewage system, in order to reduce the chemical oxygen demand (COD) of the effluent.

Problem to be Solved by the Invention

A problem with the treatment method disclosed by DE-A-3635219 is that, while the COD of the thiosulphate-containing photographic effluent is reduced, transition metal contaminants are added to the effluent as catalyst which are subsequently discharged into the environment. This is clearly undesirable for environmental reasons, and is also an inefficient use of the transition metals. It is an object of the present invention to provide an improved treatment method for waste effluent, and in particular photographic effluent, containing reduced species, which does not involve the discharge of transition metal species to the environment and in which the transition metal species can be reused.

Summary of the Invention

According to one aspect of the present invention therefore there is provided a method of treating waste effluent containing reduced species, by oxidation with hydrogen peroxide, or a compound capable of releasing hydrogen peroxide, in the presence of a catalyst therefor, characterized in that said catalyst is immobilized on a substrate therefor.

The reduced species may generally be a sulphur-oxygen species, typically as thiosulphate or sulphite.

Said catalyst may be selected from molybdate, tungstate, chromate and vanadate, although tungstate and especially molybdate are preferred.

Typically the substrate will constitute a porous mass which permits permeation of the photographic effluent into its interstices, thereby presenting a large surface area of catalyst to the effluent. Said substrate may be anionic, and in a particular aspect of the present invention the substrate comprises an anion exchange material. In some embodiments, a mixture of anion and cation exchange materials may be used. The material may typically be a polymeric resin, especially a polyacrylic or polystyrene resin, or clay or zeolite-type material.

In another aspect, the present invention contemplates a photographic development process in which effluent from one or more of the development steps are treated continuously or batch-wise by the method according to the present invention. Usually, the effluent from each stage will be combined and treated together. Said development process may be a redox-amplification (RX) process which is performed upon a photographic element containing a reduced silver lay-down density as compared with conventional silver halide photographic materials. As will be well known to a person skilled in the art, hydrogen peroxide is employed in the development step of an RX process, in which case it may be unnecessary to add additional peroxide to the photographic effluent. Of course, additional peroxide may be added to the effluent if required.

The above method is therefore particularly appropriate for use with dilute solutions containing fixer, for example those with less than about 20g

thiosulphate (based on ammonium thiosulphate), and hence is suitable for RX developed images.

5 However it has been found that with those fixer solutions wherein
the amount of sulphur-oxygen species is high, i.e. conventional solutions such as
those used in x-ray and graphic arts, and wherein sufficient peroxide is added to
effect total oxidation to sulphate, the pH may become very acid (even as low as
pH 1) which would be corrosive and unsuitable for discharge directly to sewers.
10 The effluent could be subsequently neutralized by adding an alkali or base to the
treated liquid in an amount appropriate to render the pH suitable for disposal,
generally around pH 5-9, but this would require a separate process. Adding an
alkali metal hydroxide or carbonate having a high pH to the added peroxide
solution however causes the peroxide to decompose rapidly and therefore it
cannot be stored therewith.

15 In accordance with a further aspect of the invention the above
problem has been overcome by the addition to the peroxide solution of a soluble
alkali whose conjugate acid has a pKa of < 8.5, such as for example a soluble
bicarbonate, alkanoate, e.g. acetate, or dihydrogen phosphate, in an amount
20 approximately equivalent to the acid released in the decomposition reaction. Only
very slight decomposition of peroxide over a period of several weeks results.

 The soluble alkali is an alkali metal or ammonium salt, preferably a
bicarbonate and more preferably the highly soluble potassium bicarbonate since
25 the solubility of the alkali limits the amount of the acid that can be neutralized. At
low pH the bicarbonate is converted to carbon dioxide which escapes from the
solution thereby not adding to the salt content, which would be the case for other
salts except for hydroxide or carbonate. Potassium bicarbonate is also the least
polluting alkali, as it has no biological oxygen demand (BOD₅), and is also the
30 cheapest, making it particularly suitable for practice in the present invention.

 The pH of the effluent is raised by the above process but cannot exceed
the natural pH of the alkali (about pH 9) so if there is a pump fault anywhere in the
apparatus then the pH of the effluent can never go too high. Moreover provided an
35 excess of alkali is added the actual amount required is not critical.

Conveniently the peroxide/alkali mixture may be stored as a combined oxidation/neutralizing solution which can then be pumped through a single pump and thence together with the fixer solution over the immobilized substrate as described hereinafter.

Compounds capable of releasing hydrogen peroxide include metal peroxides; compounds which include hydrogen peroxide in their crystal structure such as sodium percarbonate; other peroxy compounds such as sodium perborate and persulphate; or soluble organic peroxide, such as butyl peroxide or benzyl peroxide. The peroxide is added in an amount sufficient to cause oxidation of a substantial proportion of the reduced species and is conveniently hydrogen peroxide itself.

Where the photographic effluent is treated batch-wise, the catalyst may be disposed within a receptacle which is equipped with an inlet for introducing photographic effluent from the development process and an outlet for discharging treated effluent to waste. The outlet will be fitted with selectively operable closing means for closing the outlet during conduction of the treatment method, typically a valve.

Alternatively, the treatment method may be performed continuously on effluent delivered from the photographic development process. In another aspect of the present invention therefore the catalyst/substrate may be packed in a conduit which is arranged to receive photographic effluent in one end, and to deliver the treated photographic effluent from the other end. The flow rate of photographic effluent through the conduit will be adjusted such that the average residence time of effluent within the conduit is sufficient to oxidize a substantial proportion of the reduced sulphur-oxygen species contained in the effluent.

In the method wherein alkali is added to the peroxide in storage, the experimental set-up is as described hereinafter.

Where thiosulphate-containing effluents from the fixing stage are treated in accordance with the present invention, silver that has been complexed

during fixing may be precipitated in the treatment stage. In some embodiments therefore silver-bearing precipitate may be separated from the treated photographic effluent before the effluent is discharged. For this, filtering or centrifuging means may be employed.

5

Advantageous Effect of the Invention

According to the present invention therefore a method of treating waste effluent containing reduced species, preferably containing sulphur-oxygen species, is provided which does not involve discharging transition metal species into a sewage system. This is advantageous from the environmental perspective, and also means that the transition metal catalysts can be reused which is more efficient in these materials as compared with the prior art processes.

Unexpectedly, the present applicants have found that by supporting the transition metal catalysts on a substrate therefor, substantially less catalyst can be used as compared with the prior art processes without impairing the efficiency of the treatment reaction. For example, the treatment method of the present invention can be performed effectively using less than 1/100 parts by weight catalyst, and typically less than 1/250 parts by weight of sulphur-oxygen species. It was also found surprisingly that use of a supported catalyst in accordance with the present invention appeared to cause or allow more complete destruction of sulphur-oxygen species to sulphate as compared with prior art processes using unsupported catalyst.

A method is further provided wherein addition of alkali, such as potassium bicarbonate, to the peroxide solution in the case of effluents with high fixer concentration enables the pH of the treated effluent to be maintained at environmentally acceptable levels.

Brief Description of the Drawings

Figures 1 to 4 of the accompanying drawings are graphs of pH verses time for reactions of photographic effluent with peroxide in the presence of a supported catalyst.

Figure 1 is a pH versus time graph for effluent reaction with peroxide with resin, without resin and a control without molybdate.

Figure 2 shows the use of a number of Dowex 1x8 series resins, and also a control.

5 Figure 3 shows the use of Dowex 1x2 series resins, and also a control.

 Figure 4 shows the use of a Duolite 6113 mixed-bed resin, and a control.

10 Figure 5 shows the holding tank for performing the treatment method of the present invention.

 Figure 6 shows a conduit in accordance with the present invention for performing the treatment method.

 Figure 7 shows an experimental set-up to demonstrate the use of a storable peroxide/alkali combination in a process according to the invention.

20 The invention will now be described with reference to the following Examples which are not to be construed as limiting in any way.

Detailed Description of the Invention

EXAMPLE 1

25 A "mock" effluent from a low silver RX color process was made up from the following solutions:

Developer:

	1-hydroxyethylidene-1,1'-	
	-diphosphonic acid	0.6g
30	Diethyltriaminepentaacetic acid	2.0g
	Dipotassium hydrogen phosphate	40.0g
	Hydroxylamine sulphate	0.5g
	CD3	4.5g
	Hydrogen Peroxide (30%)	2ml
35	Water to	1 litre

pH adjusted to 11.7

[CD3 = 4-amino-3-methyl-N-ethyl-(2-methanesulfonamido-ethyl)aniline
sesquisulfate hydrate.]

5

A pseudo seasoned fix/stop was made up with the following
constitution:

	Sodium thiosulphate pentahydrate	20g
	Sodium meta bisulphite	30g
10	Sodium acetate	40g
	Silver chloride	1.2g
	Water to	1 litre

The pH of the fix/stop measured at 25C was adjusted to 6.3 with
sodium hydroxide.

15

Effluent = 6 parts developer + 5 parts fix/stop + 20 parts water.

Molybdate ions were first adsorbed on an anionic resin using the
following method. 10g of Amberlite® IRA-400 exchange resin (manufactured by
20 Rohm and Haas) was stirred gently in 50ml demineralised water. To this was
added 2ml 1% ammonium molybdate solution. This mixture was stirred for 10
minutes and then the liquid was decanted from the resin and the resin was washed
with 3 x 50ml changes of demineralised water. The drain resin was used for the
following experiments.

25

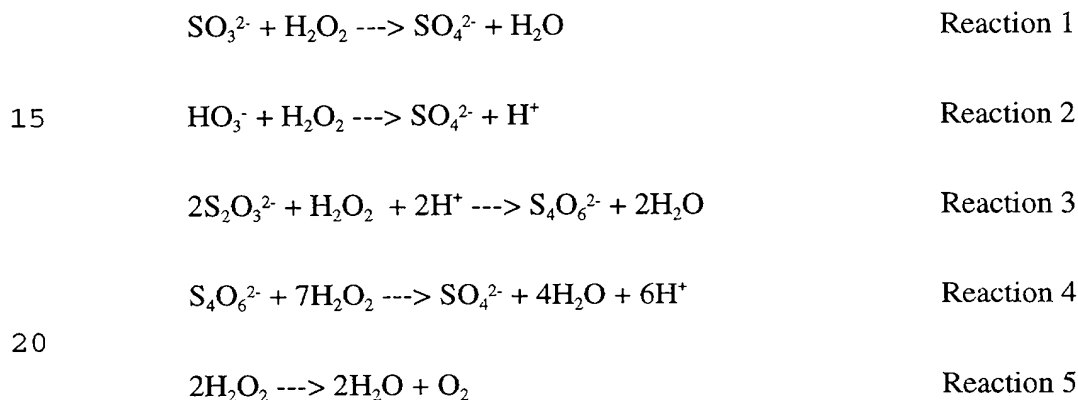
100ml of effluent at room temperature (22C) were placed in a
150ml beaker on a magnetic stirrer and stirred gently. A glass pH electrode and
calomel reference electrode were placed in the solution and connected to a
Radiometer pH meter, the meter having being calibrated previously with reference
30 pH buffers. The pH was recorded. 30ml of 3% hydrogen peroxide were quickly
added and the pH was recorded after 10sec and then as felt appropriate until the
pH stayed constant. At the end of the run the treated liquid was removed and
analyzed for molybdenum by atomic adsorption spectroscopy. To demonstrate
the invention the experiment was repeated with the 10g treated resin added. At
35 the end of the run the resin was filtered off before the treated liquid was sent for

molybdenum analysis as before. The experiment was again repeated but this time with 2ml 1% ammonium molybdate added directly to the mixture of effluent and peroxide with no resin present.

- 5 The levels of molybdenum in the treated effluent are shown in the following table:

Effluent treatment	MO in Final Effluent PPM
Peroxide only	< 0.05
Peroxide + molybdate adsorbed on resin	0.55
Peroxide + molybdate	86

- 10 The destruction of sulphite and thiosulphate in the "mock" photographic effluent was monitored by observing the pH change with time. The reactions taking place in the mixture are as follows:



- 25 The pH of the mixture starts at around 7. At this pH almost all the sulphite in the mixture will be protonated as the bisulphite and of the first two reactions, reaction 2 will be the more important, so as the reaction takes place the pH should fall as protons are liberated. Reaction 3 must take place before reaction 4. Reaction 3 causes the pH to rise as protons are used and the reaction 4 takes over and the pH falls again. Thus the pH can be used to follow the reaction. Reaction 5, the peroxide decomposition reaction does not change the
- 30 pH.

The pH versus time curves for these runs are shown in Figure 1. The initial pH in all runs fell rapidly leaving the thiosulphate to react. The oxidation of thiosulphate in the run containing the resin with molybdate adsorbed was about three time faster than the run without a catalyst. The final pH was also lower suggesting that the reaction had proceeded further. Although the thiosulphate was removed faster and more completely by the run containing the equivalent amount of unadsorbed molybdate, the molybdenum in the effluent was about 150 times greater. This experiment shows that molybdate adsorbed on a resin is an efficient catalyst for the oxidation of photographic effluent with very little molybdenum entering the waste stream.

EXAMPLE 2

Example 1 was repeated with a series of Dowex 1X8 anionic ion-exchange resins. This series has resins of different bead sizes, the last number in the resin's name being the approximate mesh size of the beads, 50 being the largest and 400 the smallest. The 100 mesh beads were found to provide the best catalysis for the reaction, which was surprising as it would have been expected that the smaller bead with the greater surface area would be the most effective. The resulting pH curves are shown with a control, containing no molybdate, as Figure 2.

EXAMPLE 3

Example 1 was repeated with two Dowex 1X2 anionic ion-exchange resins, having a different amount of cross-linking compared to the 1X8 series. The resulting pH curves are shown with a control, containing no molybdate, as Figure 3. The reaction rates were similar to those in Example 2, but there was less difference between the 100 and 400 mesh size resins.

EXAMPLE 4

Example 1 was repeated with a Duolite 6113 resin, a mixed bed resin comprising a mixture of anionic and cationic resins. The resulting pH curves are shown with a control, containing no molybdate, as Figure 4. The mixed bed resin + molybdate showed some increased reaction rate over the control but was not as good as the pure anionic exchange resins.

In practice, the treatment method according to the present invention may be performed batch-wise in a holding tank apparatus (10) as illustrated in Figure 5 or using a continuous feed from a photographic development process, in which case a conduit apparatus (20) of the kind
5 illustrated in Figure 6 may be employed.

The holding tank (17) is equipped with an inlet (11) for receiving photographic effluents batch-wise from a photographic development process. The effluents from the various stages of the process may be treated separately, or may
10 preferably be combined and treated together. The holding tank (17) is also fitted with an outlet (12) which is provided with a manually operable valve (13). The outlet (12) is arranged for discharging treated effluents from the holding tank (17) to waste, e.g. in a public utility sewage system, via a separator (14) for separating precipitated silver species from the treated effluents.

15 The holding tank (17) contains a bed (15) of anion exchange resin (16). Alternatively a mixed bed of cation and anion exchange resins may be used. The exchange resin(s) (16) is prior-treated with a solution of a chromate, vanadate or preferably tungstate or molybdate salt as hereinbefore described, so as to
20 immobilize the transition metal oxyanion on the exchange resin(s).

In service, photographic effluents from the development process are introduced batch-wise to the holding tank (17) through inlet (11) with valve (13) in the closed position. If necessary, peroxide, in the form of hydrogen
25 peroxide, or a metal or organic peroxide capable of releasing hydrogen peroxide on contact with water, may be introduced to the holding tank (17) at this stage. Where the treatment is carried out on effluents from an RX development process however, the effluents may themselves already contain sufficient hydrogen
peroxide.

30 The effluents and peroxide are allowed to stand in the holding tank (17) in contact with the ion exchange resin (16) for a predetermined period of time sufficient to allow substantially complete reaction of the hydrogen peroxide with sulphite and thiosulphate anions in the effluents, for example, for a period of less

than 1 hour. This reaction is catalyzed by the transition metal oxyanions supported on the exchange resin beads (16), and forms sulphate anions.

5 The valve (13) is then opened, allowing the effluents to be discharged in the holding tank (17) through the outlet (12). Any precipitated silver species are removed by the separator (14), which may be a filter or centrifuge, and the effluents can then be discharged to waste.

10 Alternatively, the photographic effluents can be introduced continuously from the photographic development process to the conduit apparatus (20) as shown in Figure 6. In this embodiment, the conduit (27) is generally U-shaped, but different configurations may be employed as desired. The conduit (27) comprises an inlet (21) and an outlet (22) and is packed as shown in Figure 6 with a porous bed of ion exchange resin (26) as previously described. The outlet
15 (22) is equipped with a manually adjustable valve (23) which is operated to control the flow rate of effluents through the conduit (27), such that the average residence time of the effluent in the conduit (27) is sufficient to allow substantially complete destruction of sulphite and thiosulphate to sulphate, for example, preferably less than 1 hour. The effluent is then discharged to waste via a
20 separator (24) as hereinbefore described.

EXAMPLE 5

A fixer solution comprising seasoned fixer taken from a graphic arts processor was analyzed and found to contain:

25

ammonium thiosulphate	142g/l
sodium sulphite	12g/l
silver (complexed)	16.2g/l

30

The formula of a hydrogen peroxide/potassium bicarbonate treating solution was as follows:

30% hydrogen peroxide	250ml
potassium bicarbonate	50g
35 water to	1 litre

The experiment was run with and without bicarbonate in the solution, the final pH being measured in both cases. In the run without bicarbonate the pH of the treated effluent was at an environmentally unacceptable pH of 2.1. When bicarbonate was present together with the peroxide the pH of the treated effluent was satisfactory at 6.5.

To test the efficacy of the peroxide/bicarbonate solution in storage, the combined solution was kept for a period of 4 weeks under normal conditions and the above experiment repeated. The pH was found to be 6.5 as before. The level of peroxide in the combined solution was analyzed by standard iodide titration and found to have decreased by only 0.8%.

The experiment was repeated using however a peroxide/-sodium hydroxide solution containing 20g/l sodium hydroxide and having the same neutralizing power as the above. After 4 weeks the solution had completely decomposed and no peroxide was detected.

The experimental set-up for the above experiment was as shown in Figure 7. A peroxide/bicarbonate solution (30) having the quantities above and the seasoned fixer (40) described above were introduced into vessels (50) and (60) respectively. Molybdate ions were adsorbed onto an anionic resin (140) as described in Example 1 except that the resin used was Amberlite ® IRA-458 exchange resin (manufactured by Rohm and Haas), which is a polyacrylate resin. In an additional experiment the alternative polystyrene anionic exchange resin Amberlite ® IRA-68 was used.

The 'mock' effluent containing fixer solution (40) was then pumped via pump (80) to the holding tank (180), which initially only contained water, within overflow vessel (110) on a magnetic stirrer (120) and the solution (90) being treated was stirred gently. A glass pH electrode and calomel reference electrode were placed in the solution and connected to a Radiometer pH meter, as described in Example 1, for measuring the pH at predetermined intervals. The solution (90) in the holding tank (180) was then pumped, via pump (130), to combine with the peroxide/bicarbonate solution (30), pumped via pump (70), to

pass over the resin bed, equipped with filters (150,160) at either end and containing resin (140). After passing through the bed the treated effluent (190) was introduced back into the holding tank (180) and allowed to form an overflow (100) into the vessel (110).

CLAIMS

1. A method of treating waste effluent containing reduced species
by oxidation with hydrogen peroxide, or a compound capable of releasing
5 hydrogen peroxide, in the presence of a catalyst therefor, characterized in that said
catalyst is immobilized on a substrate therefor.
2. A method as claimed in claim 1 characterized in that the effluent
is photographic effluent.
10
3. A method as claimed in either of the preceding claims
characterized in that the reduced species are sulphur-oxygen species.
4. A method as claimed in claim 3 characterized in that the
15 sulphur-oxygen species are thiosulphate or sulphite.
5. A method of treating waste effluent as claimed in any one of the
preceding claims characterized in that the catalyst is selected from a molybdate,
tungstate, chromate and vanadate.
20
6. A method as claimed in claim 5 characterized in that the catalyst
is a molybdate.
7. A method as in claim 1 characterized in that the substrate
25 constitutes a porous mass which permits permeation of the waste effluent into its
interstices, thereby presenting a large surface area of catalyst to the effluent.
8. A method as in claim 1 characterized in that the substrate is an
ion exchange material.
30
9. A method as in claim 1 8 characterized in that the substrate
comprises an anion exchange material.
10. A method as in claim 1 characterized in that the effluent is
35 from a process with a redox-amplifier developer.

11. A method as in claim 1 characterized in that the effluent is from a process wherein the fixer contains an amount of sulphur-oxygen species greater than about 20g of thiosulphate (based on ammonium thiosulphate).

5

12. A method as in claim 1 characterized in that hydrogen peroxide, or a compound capable of releasing hydrogen peroxide, is combined with a soluble alkali whose conjugate acid has a pKa of < 8.5, prior to reaction with the effluent, to reduce the final pH of the effluent to about 5 to 9.

10

13. A method as in claim 12 characterized in that the alkali is a soluble bicarbonate, alkanoate or dihydrogen phosphate.

15

14. A method as in claim 13 characterized in that the alkali is potassium bicarbonate.

15. Holding tank apparatus (10) for treating waste effluents, which holding tank apparatus (10) comprises a receptacle (17) containing a catalyst, which catalyst is adapted for catalyzing the oxidation of reduced species in waste effluents by hydrogen peroxide, or a compound capable of releasing hydrogen peroxide, and which catalyst is immobilized on a substrate (16) therefor in the receptacle (17), an inlet (11) for introducing effluent from a development process to the receptacle (17), and an outlet (12) fitted with selectively operable closing means (13).

20
25

16. Holding tank apparatus (10) as claimed in claim 15 characterized in that the waste effluent is as claimed in any one of claims 2 to 4, 11 and 12.

30

17. Conduit apparatus (20) for treating waste effluents, which conduit apparatus (20) comprises a conduit (27) containing a catalyst, which catalyst is adapted for catalyzing the oxidation of reduced species in waste effluents by hydrogen peroxide, or a compound capable of releasing hydrogen peroxide, and which catalyst is immobilized on a substrate (26) therefor, an inlet

35

(21) for introducing waste effluents to the conduit (27), and an outlet (22); whereby in use, waste effluents are supplied continuously to the conduit (27) at a volume throughput to achieve substantially complete oxidation of the reduced species.

5

18. Conduit apparatus (20) as claimed in claim 17, characterized in that the substrate (26) is porous and is packed in the conduit (27).

19. Conduit apparatus (20) as claimed in either claim 17 or 18
10 characterized in that the waste effluent is as claimed in any one of claims 2 to 4, 10 and 11.

20. Apparatus for treating waste effluents (40) in a continuous
manner as claimed in any one of claims 18 to 20 characterized by including a
15 pump (130) for pumping waste effluent from a holding tank (180), a pump (70) for pumping hydrogen peroxide, or a compound capable of releasing hydrogen peroxide, or its combination with a soluble alkali whose conjugate acid has a pKa of < 8.5, for mixing with the waste effluent prior to passing over the catalyst immobilized on the substrate (140).

20

ABSTRACT

This invention provides a method of treating waste effluent, particularly photographic effluent, containing reduced species such as thiosulphate or sulphite, by oxidation with hydrogen peroxide or a compound capable of releasing hydrogen peroxide, in the presence of a catalyst therefor. The invention is characterized in that the catalyst is immobilized on a substrate. The catalyst may be selected from chromate, vanadate and preferable molybdate or tungstate and the substrate may comprise an ion exchange material, especially an anion exchange material. The invention further provides a holding tank apparatus (10) or a conduit apparatus (20) for carrying out this method.

The invention is particularly appropriate for use with fixer from redox-amplification processes. With conventional solutions with higher concentrations of fixer a soluble alkali, especially potassium bicarbonate, may be combined with the oxidizing agent and stored for at least 4 weeks without decomposition, whereby the pH of the final effluent is rendered environmentally acceptable.

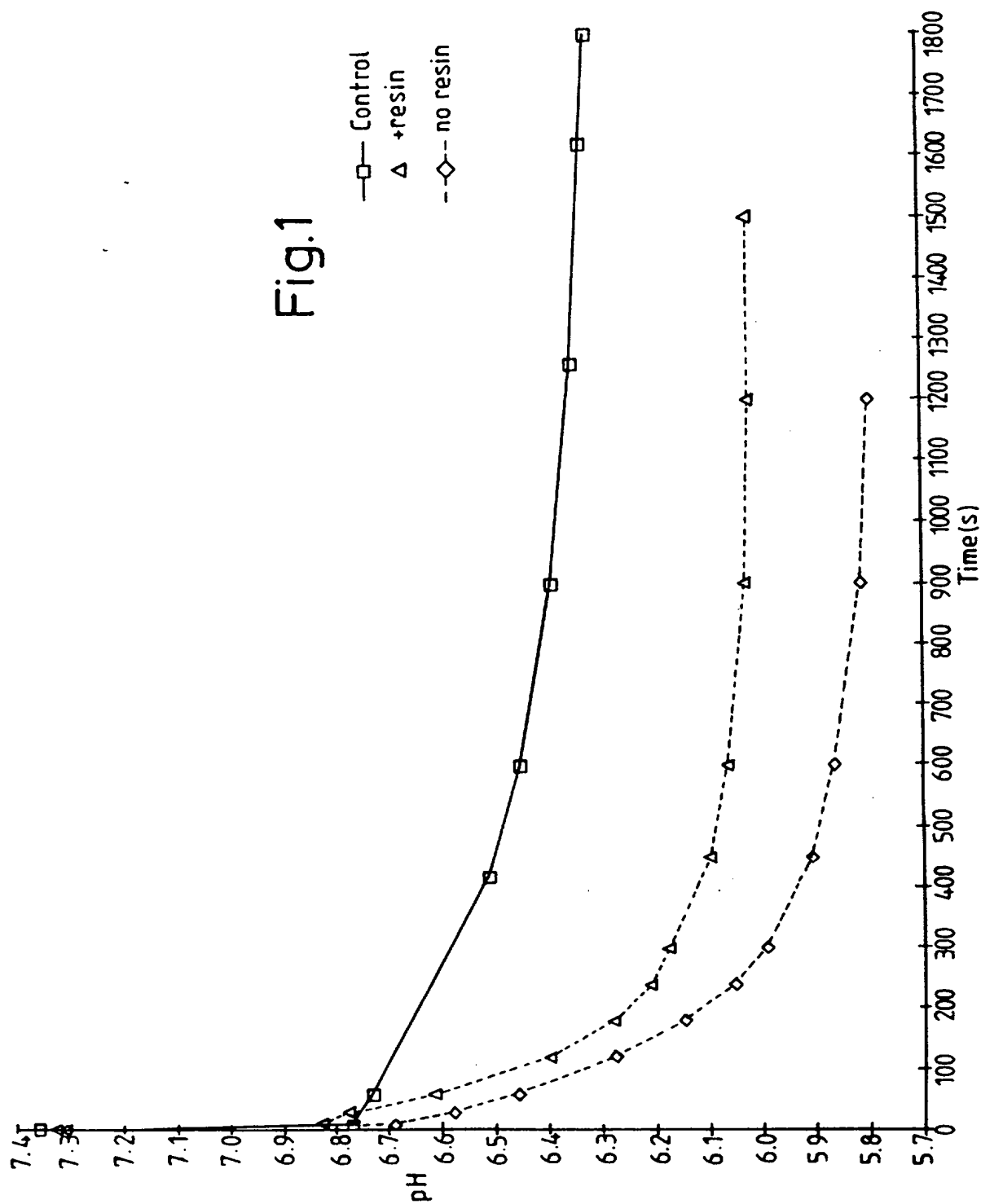


Fig.2

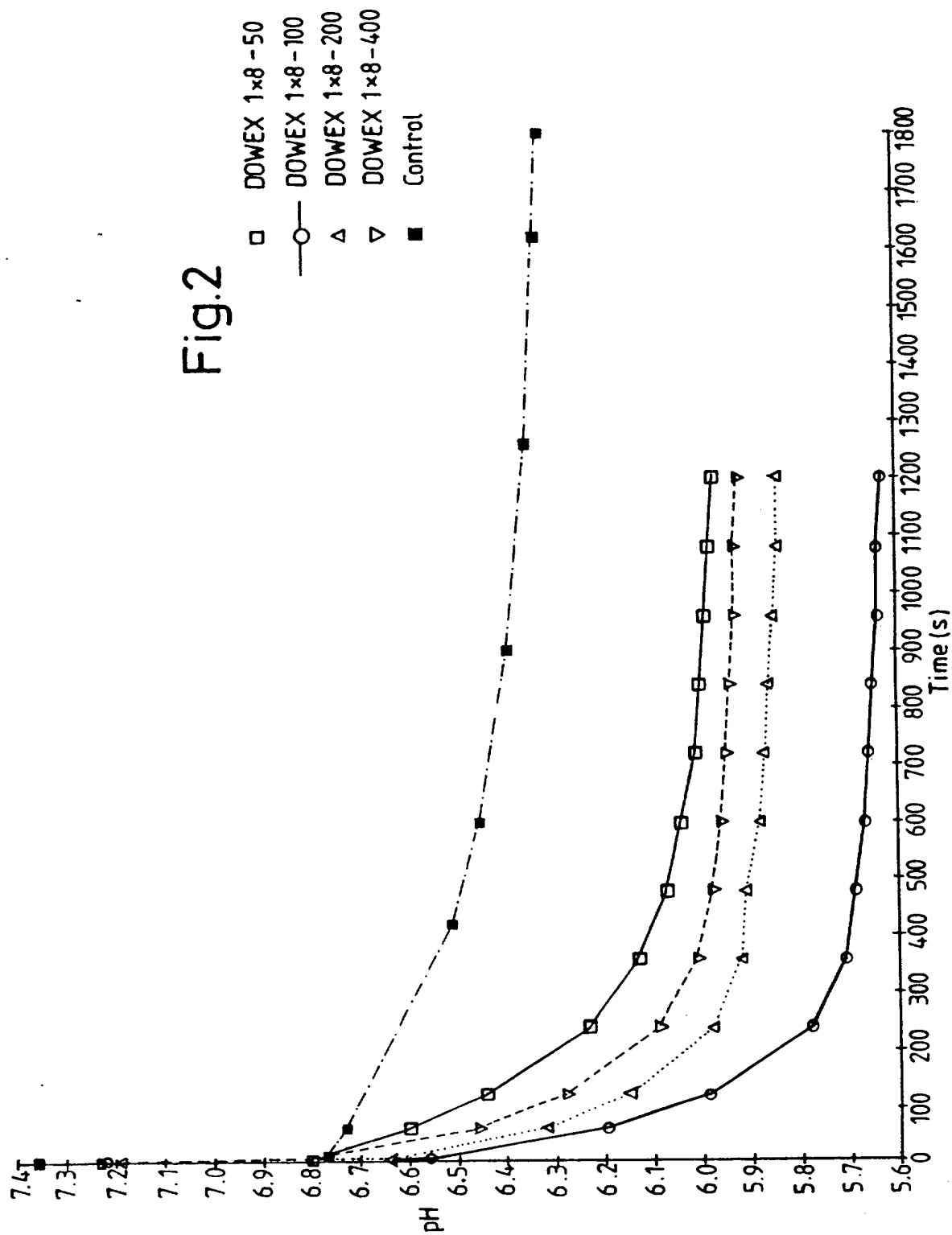
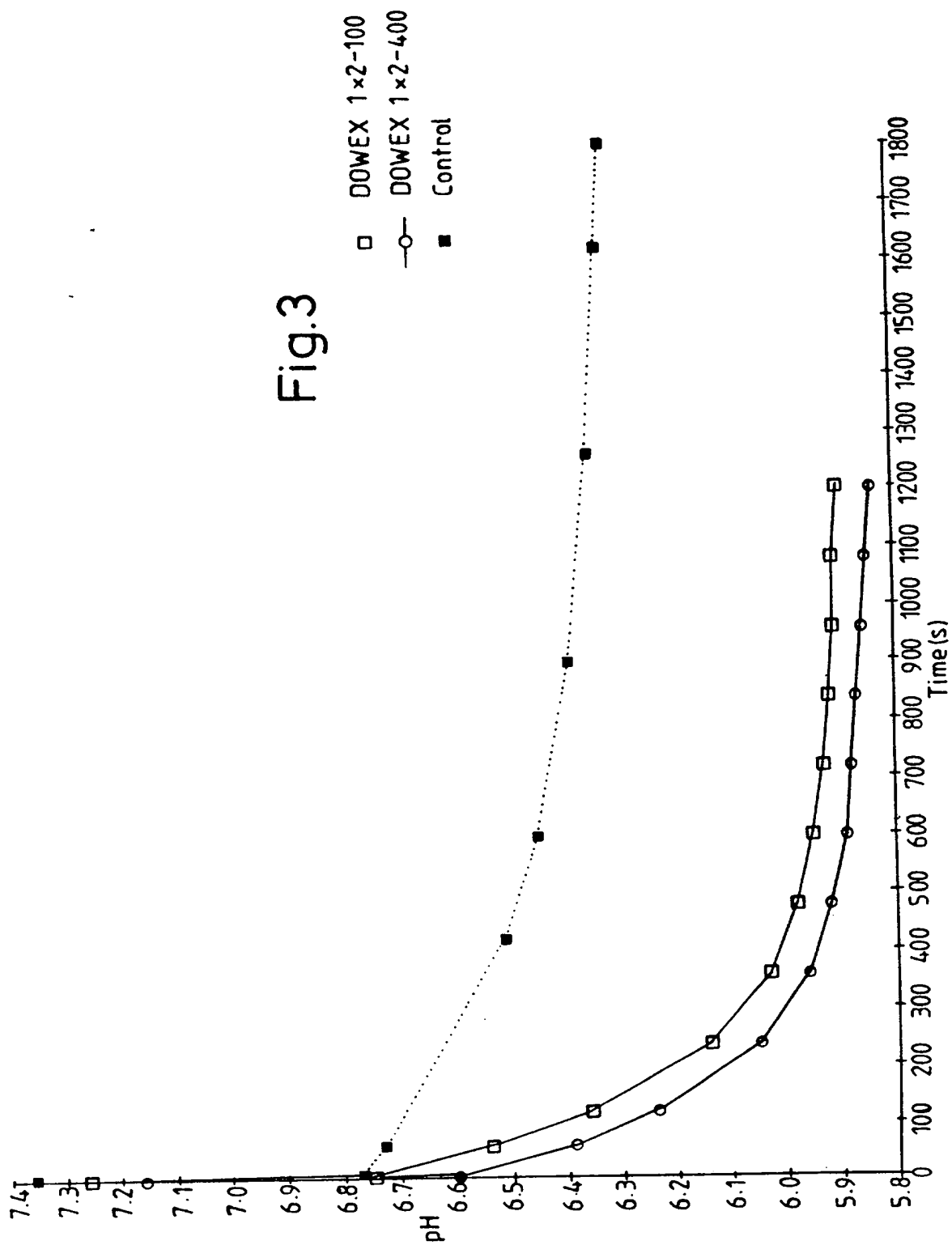
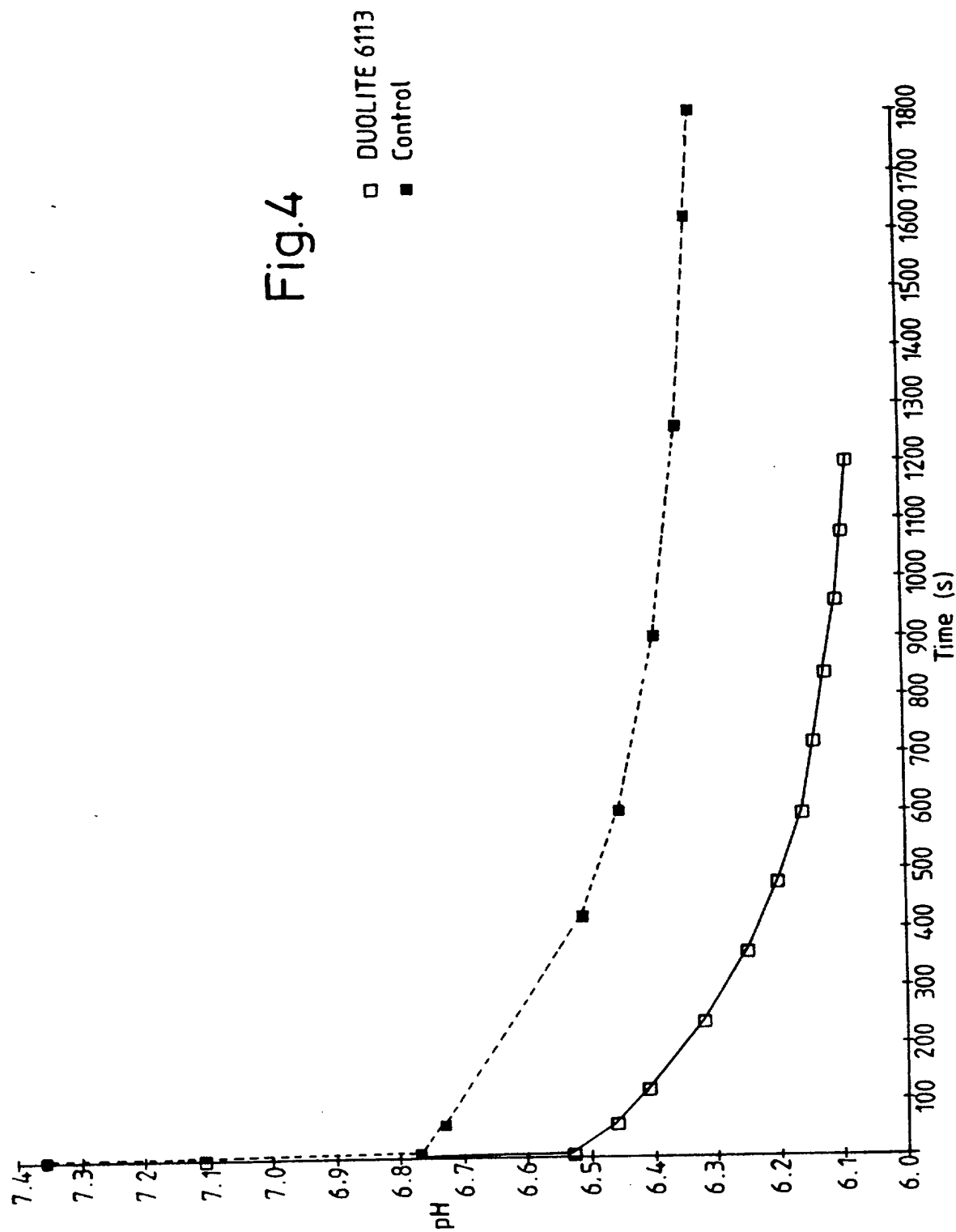
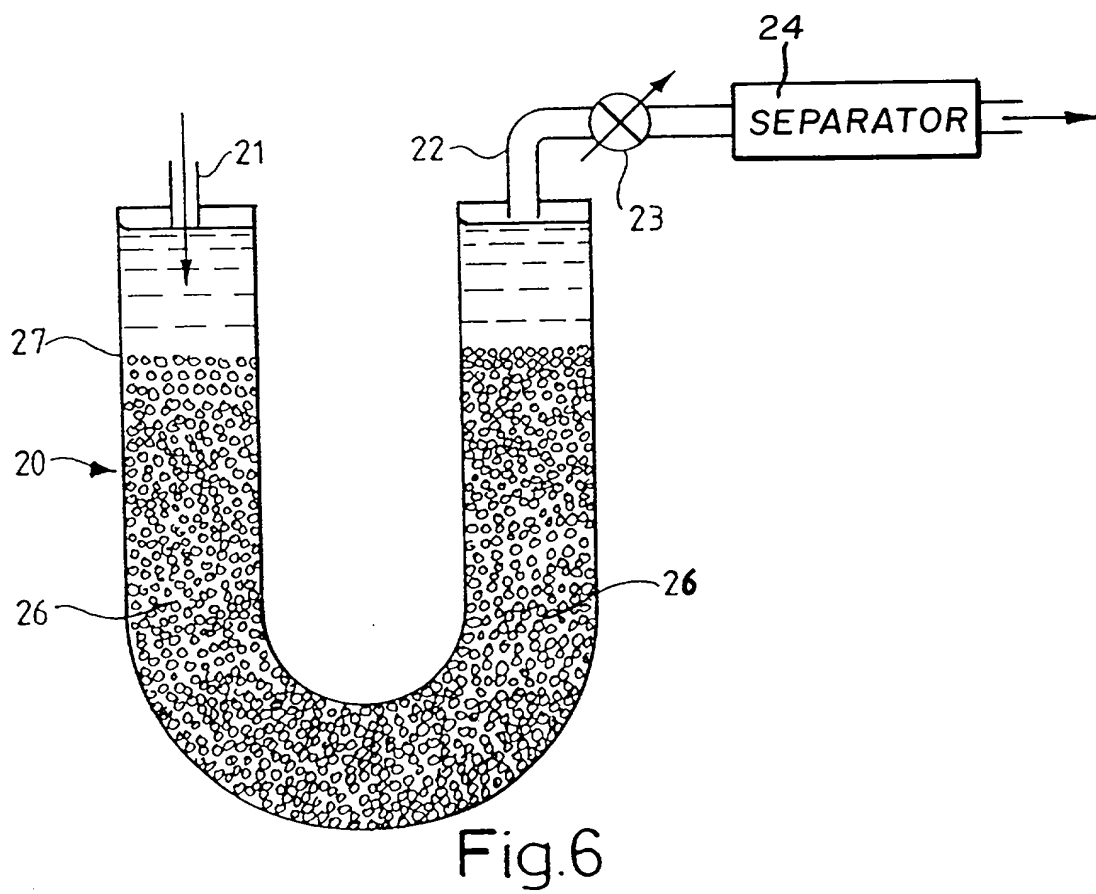
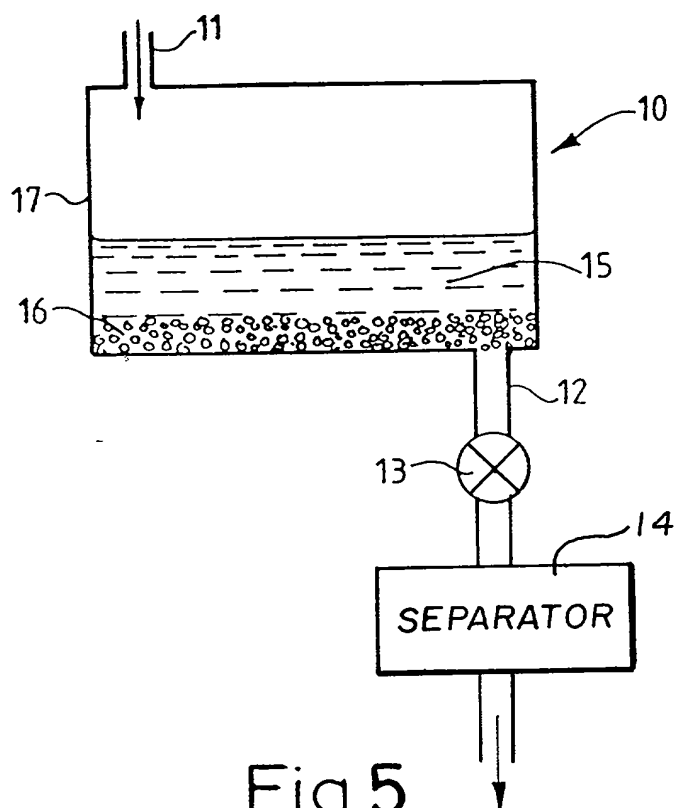


Fig.3







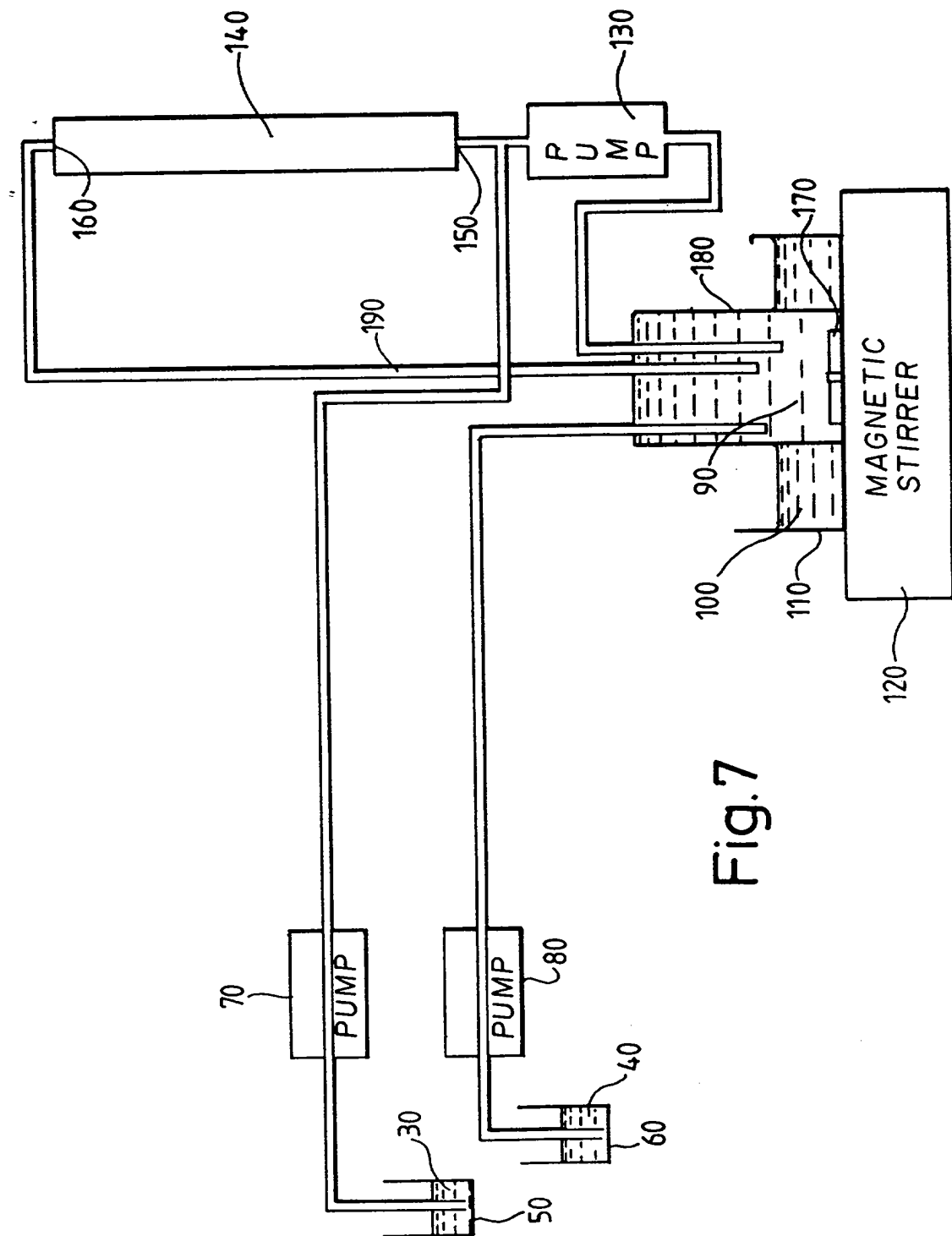


Fig.7

Combined Declaration For Patent Application and Power of Attorney**ATTORNEY DOCKET
71442JRE**

As below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

METHOD OF TREATING WASTE EFFLUENT

The specification of which (check only one item below):

☒ is attached hereto.☐ was filed as United States Application Serial No. on and
was amended on (if applicable).☐ was filed as PCT international application Number on and was amended under PCT Article 19 on (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose to the U.S. Patent & Trademark Office all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign applications(s) for patent or inventor's certificate or any PCT international application(s) designating a least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) of which priority is claimed:

PRIOR FOREIGN/PCT APPLICATION(S) AND ANY PRIORITY CLAIMS UNDER 35 U.S.C. 119:

COUNTRY (if PCT, indicate PCT)	APPLICATION NUMBER	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 USC §119			
UNITED KINGDOM	9601915.3	JANUARY 31, 1996	<input checked="" type="checkbox"/>	YES	<input type="checkbox"/>	NO
			<input type="checkbox"/>	YES	<input type="checkbox"/>	NO
			<input type="checkbox"/>	YES	<input type="checkbox"/>	NO

I hereby claim the benefit under Title 35, United States Code, 119 §(e) of any United States provisional application(s) listed below:

PRIOR PROVISIONAL APPLICATION(S) AND ANY PRIORITY CLAIMS UNDER 35 U.S.C. §119 (e):

PROVISIONAL APPLICATION NUMBER	FILING DATE

I hereby claim the benefit under Title 35, United States Code, §120 of any prior United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior applications(s) in the manner provided by the first paragraph of Title 35, §112, I acknowledge the duty to disclose to the U.S. Patent & Trademark Office all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations §1.56, which became available between the filing date of the prior application(s) and the national or PCT international filing date of this application:

PRIOR US APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS DESIGNATING THE U.S. FOR BENEFIT UNDER 35 USC §120:

U.S. APPLICATIONS			STATUS (Check one)		
U.S. APPLICATION NUMBER	U.S. FILING DATE		PATENTED	PENDING	ABANDONED
PCT APPLICATIONS DESIGNATING THE U.S.					
PCT APPLICATION NO.	PCT FILING DATE	U.S. SERIAL NUMBERS ASSIGNED (if any)			

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith (*List name and registration number*)

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

SIGNATURE OF INVENTOR 201	SIGNATURE OF INVENTOR 202	SIGNATURE OF INVENTOR 203
<i>John Richard Fyson</i>		
DATE	DATE	DATE
18 December 1996		
SIGNATURE OF INVENTOR 204	SIGNATURE OF INVENTOR 205	SIGNATURE OF INVENTOR 206
DATE	DATE	DATE